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3151 CTGCTTCGAA GATGGCACCT TTTTGCTGTG TTCTCA (SEQ ID NO: 1)

FEATURES:

5'UTR: 1-14 Start Codon: 15 Stop Codon: 1188 3'UTR:

Homologous proteins:

Top 10 BLAST Hits:	Score	E
CRA 335001098638983 /altid=gi 11321561 /def=ref NP_000604.1 he	681	0.0
CRA 18000004928118 /altid=gi 386789 /def=gb AAA52704.1 (J03048	679	0.0
CRA 18000005034645 /altid=gi 1335098 /def=emb CAA26382.1 (X025	634	0.0
CRA 18000004885233 /altid=gi 1708184 /def=sp P20058 HEMO_RABIT	519	e-146
CRA 18000004905757 /altid=gi 1070649 /def=pir OQRB hemopexin p	513	e - 144
CRA 84000015361878 /altid=gi 13641048 /def=ref XP_011963.2 hem	504	e-141
CRA 18000004936853 /altid=gi 123036 /def=sp P20059 HEMO_RAT_HEM	466	e-130
CRA 18000004950055 /ditid=gi 1708183 /def=sp P50828 HEMO_PIG_HE	459	e-128
CRA 18000005011238 /altid=gi 1087020 /def=gb AAA82488.1 hepato	436	e-121
CRA 18000005041763 /altid=gi 1311343 /def=pdb 1HXN Heme Mol	408	e-113
Blast hits to dbEST:	Score	F
0.60	1360	0.0
gi 12798347 /dataset=dbest /taxon=960	1344	0.0
gi 12914625 /dataset=dbest /taxon=960	973	0.0
gi 6360478 /dataset=dbest /taxon=9606	967	0.0
gi 9866417 /dataset=dbest /taxon=960	839	0.0
α ; 112798348 /dataset=dhest /taxon=960	0.00	0.0

Expression Information:

Tissue source of BLAST dbEST hits:

gi|12798347 Fetal brain

gi|12914625 brain neuroblastoma cells

gi|12798348 /dataset=dbest /taxon=960...

gi|6360478 liver

gi|9866417 non cancerous liver tissue

gi|12798348 Fetal brain

Tissue source of cDNA clone:

Fetal liver

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  51 SDGWSFDATT LDDNGTMLFF KGEFVWKSHK WDRELISERW KNFPSPVDAA
 101 FRQGHNSVFL IKGDKVWVYP PEKKEKGYPK LLQDEFPGIP SPLDAAVECH
 151 RGECQAEGVL FFQGHGHRNG TGHGNSTHHG PEYMRCSPHL VLSALTSDNH
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 301 LHIMAGRRLW WLDLKSGAQA TWTELPWPHE KVDGALCMEK SLGPNSCSAN
 351 GPGLYLIHGP NLYCYSDVEK LNAAKALPQP QNVTSLLGCT H (SEQ ID NO:2)
FEATURES:
Functional domains and key regions:
Prosite results:
PDOC00001 PS00001 ASN GLYCOSYLATION
N-glycosylation site
Number of matches: 4
            64-67 NGTM
      1
                           NGTG
             169-172
                           NSTH
      3
             175-178
             382-385
                           NVTS
       4
PDOC00005 PS00005 PKC PHOSPHO_SITE
Protein kinase C phosphorylation site
Number of matches: 5
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             47-49 TER
             78-80 SHK
             87-89 SER
       3
            216-218
                           TSR
       4
             298-300
                           SSR
       5
PDOC00006 PS00006 CK2_PHOSPHO_SITE
Casein kinase II phosphorylation site
Number of matches: 9
                     TKPD
           40-43
      1
             59-62 TTLD
       2
            95-98 SPVD
       3
                           SPLD
            141-144
                           TSRD
            216-219
                           SAVD
            235-238
       7
            242-245
                           SWEE
             321-324
                           TWTE
       8
             366-369
                           SDVE
       9
PDOC00008 PS00008 MYRISTYL
N-myristoylation site
Number of matches: 5
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       1
             170-175
                          GTGHGN
       2
             201-206
                           GATYAF
       3
                           GTPHGI
       4
              279-284
                           GAQATW
             317-322
 PDOC00009 PS00009 AMIDATION
Amidation site
              305-308
                            AGRR
 PDOC00013 PS00013 PROKAR_LIPOPROTEIN
 Prokaryotic membrane lipoprotein lipid attachment site
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OPONVTSLLGC

379-389

PDOC00023 PS00024 HEMOPEXIN Hemopexin domain signature

Number of matches: 2

1 86-101 ISERWKNFPSPVDAAF

2 226-241 IAHQWPQGPSAVDAAF

Membrane spanning structure and domains:

elix	Begin	End	Score	Certainity
1	6	26	1.820	Certain
2	251	271	0.639	Putative

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BLAST Alignment to Top Hit:
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             hemopexin [Homo sapiens] /org=Homo sapiens /taxon=9606
              /dataset=nraa /length=462
             Length = 462
 Score = 681 \text{ bits } (1737), \text{ Expect = } 0.0
 Identities = 341/468 (72%), Positives = 351/468 (74%), Gaps = 83/468 (17%)
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              MARVLGAPVALGLWSLCWSLAIATPLPPTSAHGNVAEGETKPDPDVTERCSDGWSFDATT
Sbjct: 1 MARVLGAPVALGLWSLCWSLAIATPLPPTSAHGNVAEGETKPDPDVTERCSDGWSFDATT 60
Query: 61 LDDNGTMLFFKGEFVWKSHKWDRELISERWKNF------ 93
              LDDNGTMLFFKGEFVWKSHKWDRELISERWKNF
Sbjct: 61 LDDNGTMLFFKGEFVWKSHKWDRELISERWKNFPSPVDAAFRQGHNSVFLIKGDKVWVYP 120
Query: 94 -----PSPVDAAFR--QGH---NSVFLIKGDKVWVYP---PEKKEK 126
                             PSP+DAA +G V +GD+ W + KE+
Sbjct: 121 PEKKEKGYPKLLQDEFPGIPSPLDAAVECHRGECQAEGVLFFQGDREWFWDLATGTMKER 180
Query: 127 GYPK------LLQDEFPG-IPSPLDAAVECHRGECQAEGVLFFQ 163
               +P L D G +P V + C +
Sbjct: 181 SWPAVGNCSSALRWLGRYYCFQGNQFLRFDPVRGEVPPRYPRDVRDYFMPCPG----R 234
Query: 164 GHGHRNGTGHGNSTHHGPEYMRCSPHLVLSALTSDNHGATYAFSGTHYWRLDTSRDGWHS 223
              GHGHRNGTGHGNSTHHGPEYMRCSPHLVLSALTSDNHGATYAFSGTHYWRLDTSRDGWHS
Sbjct: 235 GHGHRNGTGHGNSTHHGPEYMRCSPHLVLSALTSDNHGATYAFSGTHYWRLDTSRDGWHS 294
Query: 224 WPIAHQWPQGPSAVDAAFSWEEKLYLVQGTQVYVFLTKGGYTLVSGYPKRLEKEVGTPHG 283
              WPIAHQWPQGPSAVDAAFSWEEKLYLVQGTQVYVFLTKGGYTLVSGYPKRLEKEVGTPHG
Sbjct: 295 WPIAHQWPQGPSAVDAAFSWEEKLYLVQGTQVYVFLTKGGYTLVSGYPKRLEKEVGTPHG 354
Query: 284 IILDSVDAAFICPGSSRLHIMAGRRLWWLDLKSGAQATWTELPWPHEKVDGALCMEKSLG 343
              IILDSVDAAFICPGSSRLHIMAGRRLWWLDLKSGAQATWTELPWPHEKVDGALCMEKSLG
Sbjct: 355 IILDSVDAAFICPGSSRLHIMAGRRLWWLDLKSGAQATWTELPWPHEKVDGALCMEKSLG 414
Query: 344 PNSCSANGPGLYLIHGPNLYCYSDVEKLNAAKALPQPQNVTSLLGCTH 391
               PNSCSANGPGLYLIHGPNLYCYSDVEKLNAAKALPQPQNVTSLLGCTH
Sbjct: 415 PNSCSANGPGLYLIHGPNLYCYSDVEKLNAAKALPQPQNVTSLLGCTH 462 (SEQ ID NO:4)
Hmmer search results (Pfam):
 Scores for sequence family classification (score includes all domains):
Model Description
                                                                         154.0 3.1e-45 4
 PF00045 Hemopexin
                                                                          10.9
                                                                                      0.014 2
 CE00423 E00423 stromelysin_1
 Parsed for domains:
                                                             score E-value
 Model Domain seq-f seq-t hmm-f hmm-t

        PEOGRAFI
        Domination
        Seq-1
        Seq-2
        Inhun-1
        Inhun-2
        Score
        B value

        PF00045
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        0.0096

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 5901 TCAGTGGTGA GAGATGCCCC CAACTCCCCC AATGTGCTCT CACATCTCTT
 5951 TTACTTGTAT CTCCCATCCT TGACACATTT CTCCATTGTC ATCACTGTGT
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 6051 CCTCATCTCT GAGGCATATT TCTCAATCTT GTCTGTCACG GCCCAAGCCC
 6101 CTAACTTCAT CTACCTGTCT ACCATCTACT CCCATGGCTG TGCCCCCTGT
 6151 GGACCTCTCT GGGCCCCTAT GACTCCTTGT GTTCTCCTTG CTCAATGCCC
 6201 TGCTGAGCCC TCTGGCTCTC CCTTGCTCCC TGGACCTCTA TGTGTCTCTG
 6251 TACCTCCTTG CCTCCCTTTG TTCTTGCATA TCTTTCTGAG TCCTCTGGCT
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6351 ATGTCCAGAC CCCTGGGCAT AGCACTGCCT GGGGATGAGA TGTTCTCATT
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6451 CTTCACTAGC CTCTGGAGGT TTCTCCTCTG AGTAGCCAAT GGAGATACCC
6501 CTCCCTTGAC CCGTGGCATC AATTGGTAAA AGCCATCTAA TAATACCTAG
6551 GGCTGTTCTG AGTTCAGTCA GGCAGTAAAT AGTCATGCTG CACAGTTGAG
6601 AATATCCCCA AGAGGAGTGA GCAACCACAT CACATCCAAC CTGAGATATA
6651 TGTATAATTA GGACAGTGGT AAGAATATAA AATCGTGAAA ATATTTTTTT
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6801 CACAGCAGGA TTGCACACTA TAATAAGAAC ATACAGCTAA GATGAAACAC
6851 ACACCTGTAG TGAAAATACA ACATTAAACT GAGAACATAC GCCATAGTAA
6901 GAACACATAA GTATCAAGAG AACACAGC CATGGTGGGA GCCCATTGGG
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7201 GGGCGTGGTG AGTCACGCCT GTAATCCTAG CACTTTGGGA GGCTGAGGCA
7251 GGTGGATTGC TTGAGCGGCT TGAGCCTAGG AGTTTGAGAC CAGCCTGGGC
7301 AACATGGCAA AACCTCATCT CTACAAAAA TATAAAAATT AGTCGGGTGT
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7651 GTTTTGTTGT TGTTGTTGCT GTTGTTTGAG ATGGAGTCTC ACTCTGTCAC
7701 CCAGGCTGGA GTGCAATGGC GCAATCTTGG TTCACTGCAA CCTCTGCCTC
7751 CTGGGTTCAA GCGATCCTCC TGCCTCAGCC TCCCGTATAG CTGGGACTAC
7801 AGGCGCATAC CACCATGCCT GGCTAATTTT TGTATTTTTT TTTGGGTTAC
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7901 TCACCATGTT GGCCAGGCTG GTCTCGAACT CCTGACCTCA GGTGATCCGC
7951 CTGCCTCGGC CTCCCAAAGT GCTAGGATTA CAGGCATGAG CAACCACGCC
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8151 GTAAGATTTC CAGAGAGGGA ATCTGAATGG CCCAGTCCAT ATTTTCAGAC
8201 CACACCACAT TAAAGTGGTT GATTGCCAGC CTATGTATTG GCTACATTAA
8251 TGGGTTGGGA ACTCATCATT TACTTCATTG CACAAAGCAG CATAGCTCTG
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8351 CCCAACACTG TGGGAGGCCG AGGGGGGCAG ATCACTTGAG TCCAGGAGTT
8401 CTAGACCAGC CTGGGCAACA TGGTGAAATC TCATCTCTAC TAAAAATACA
8451 AAAAATTAGC CAGGTGTGGT GGCATGCACC AGTAGTCCCA GCTGTTCAGG
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8551 ACCGTGACTG TGCCTCTGCA ATCCAGCCTG GGTGACAGAT TGAGACCCTG
8601 TCTCAAAAAA CAAATAAATA AAATAAAATA AATATGGTTC CTGAGCAGGG
8651 TAATTTCAGT GGGAAACCTC CCAGGGGAGG TGGATATGTC AGTCACCGCT
8701 GTATACTCAG TACACGGCTA ATAAGAGAAC TTGTGGTAGC AGCAAGAACA
8751 CTAGGTATTT ACTCAACAAA TATTTGTTGA GCATCTGATA AGAAGTGGGC
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9201 GGGTTTTCAA CCATAAAAGG GAGTCATCAG AAAGTCTTGA GCAGGGCTGT
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9351 CCGTGCAGTT GTCCAAATTA GAGGTGATGA CCGCTTGGAC TAGGATGATA
9401 GCAGCAGAGG TGGTGAGGAA TCACCATGAT ATATTTTGGA GGTAGAGCTG
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9651 GGGAGGCTGA GGCAGGAGAA TCGCTTGAAC CTGGGAGGTG AATGTTGCAG
9701 TGAGCCGAGA TTGCACCATT GCACTCCAGC CTGGGGAACA AGAGTGAAAC
9751 TCCGTCTCTA AATAAATGAA TGAATGAATG ATATCAGTCA GAGTAGGGAA
9801 GGGAAAAGAG GCTTCAAGAA TGACTCAGCT TTCGTGGACT CAGCAACTGA
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10001 CTCAAAGGAG AGGTCAGTCA GAGCTGACGA GAACAGATTG GAAGTCATCA
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10451 TTGTTGTGAA AAGAGGAGAA GAAAAACGGG GTGCTAGCCC AGCTACTCCC
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12551 AGCCTCCTGG GTATCTGCAC CATCTAGACC AGCAAATGTC ACTGGCAAGG
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12601 AGGTTGCAGT GCTTGGTTAT TTTCTGGTCA TAAACTGGTG AAGGCTTTGG
12651 GTTCCAAATT TGCTGACAGC TGTTTAACTG GGAATTGGGC CTAGACTATA
12701 GGTAGCTATG TCTCAGACAA GGCCCTATTC CTCCACTGCC TTTACAACCC
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13501 CGTGTTCTCT TGCAGGGAAT CCTCTAGCTT GTCTCCAGGG AACTCCCAGA
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13651 AAAATCCAAG ATCAAGGCTC CAGCAGGTTC AGTGTCTGCT GAGTGCTTGT
13701 TCTGCTTCGA AGATGGCACC TTTTTGCTGT GTTCTCA (SEQ ID NO:3)
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FEATURES:

Genewise results:

Start: 2001 2001-2083 Exon: Intron: 2084-2233 2234-2292 Exon: Intron: 2293-2413 2414-2485 Exon: Intron: 2486-2665 2666-2787 Exon: Intron: 2788-4442 Exon: 4443-4596 Intron: 4597-5774 5775-5906 Exon: Intron: 5907-10934 10935-11065 Exon: Intron: 11066-11148 11149-11311 Exon: Intron: 11312-11481 11482-11738 Exon: 11739 Stop:

Sim4 results:

1987-2083, (Transcript Position: 1-97) Exon: 2234-2292, (Transcript Position: 98-156) Exon: Exon: 2414-2485, (Transcript Position: 157-228) Exon: 2666-2787, (Transcript Position: 229-350) 4443-4596, (Transcript Position: 351-504) Exon: 5775-5906, (Transcript Position: 505-636) Exon: 10935-11065, (Transcript Position: 637-767) Exon: 11149-11311, (Transcript Position: 768-930) Exon: 11482-13737, (Transcript Position: 931-3186) Exon:

CHROMOSOME MAP POSITION:

Chromosome 11

ALLELIC VARIANTS (SNPs):

T	17	-	
1)	IV	А	

Position	Major	Minor	Domain
1106	С	T	Intron
4344	A	G	Intron
7078	T	A	Intron
10841	С	G	Intron
10850	A	G	Intron
12727	G	A	Exon, 3' UTR
13164	T	G	Exon, 3' UTR
13285	\mathbf{T}	С	Exon, 3' UTR
13654	A	G	Exon, 3' UTR
13699	G	С	Exon, 3' UTR

Context:

DNA Position

1106

7078

AGGATATCGTATAAGCACAGTAGATAAAAAATGTGTGTAAATGCAGAGTGGCAGTATCTG GGGATGCACAGTCAAAAAGAGAGTACTTTTGAATGCAGGGGGACAAAGTCTGGGTATACC CTCCTGAAAAGAAGGAGAAAGGATACCCAAAGTTGCTCCAAGATGAATTTCCTGGAATCC CATCCCCACTGGATGCAGCTGTGGAATGTCACCGTGGAGAATGTCAAGCTGAAGGCGTCC TCTTCTTCCAAGGTCAGTCCAGGCTGGAATCCAAGAACCTGGAGTAGTGGTGGGTTGGTA

AGCCAGAGTGAGAACATTCAGTAGAAGTGGTGCTTCCTTTTTAAGTTCTGGACACTGTAT
TTCATTATCTATAACCGCATCTCTGTACATGGACACCTGAAATCCTTAGGGAGTGCCCGC
CAACCCCATGATGTTGGCCTTACCTGGAAACTTAGCCACTGTTTTCCACACTTGCCTTTC
TTTCAGGCACCTGCTGATTCCAGTTTCAGCCAGGGCACAGTGCCCAACATTGCTGACCAA
GTCTTGCTCTATTTCTCCTTCTCACCTGGCCTCTTCCATCTTGGCCTCTGGATGCATTCT
[C.G]

13285

GAGAACATTCAGTAGAAGTGGTGCTTCCTTTTTAAGTTCTGGACACTGTATTTCATTATC
TATAACCGCATCTCTGTACATGGACACCTGAAATCCTTAGGGAGTGCCCGCCAACCCCAT
GATGTTGGCCTTACCTGGAAACCTTAGCCACTGTTTTCCACACTTGCCTTTCTTCAGGCA
CCTGCTGATTCCAGTTTCAGCCAGGGCACAGTGCCCAACATTGCTGACCAAGTCTTGCTC
TATTTCTCCTTCTCACCTGGCCTCTTCCATCTTGGCCTCTGGATGCATTCTCCCCTCTC
[A, G]

TGACTCATTTCTGCATTCATCACTAGCCTCTTCTCTGCCTGGGCTTCTGCCAGCGGCCCT
AGAGCAACCTATGGTATTCCACAGGGACCCACTACTGGCGTCTGGACACCAGCCGGGATG
GCTGGCATAGCTGGCCCATTGCTCATCAGTGGCCCCAGGGTCCTTCAGCAGTGGATGCTG
CCTTTTCCTGGGAAGAAAAACTCTATCTGGTCCAGGTGTATTGGGGGAGAGGCTTGAG
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12727 CAAGCTGGAGAGAGAAGAAGAATGAATGGCACCATGGAGCACATTTGCAGAACACAG
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CAGCTGTTTAACTGGGAATTGGGCCTAGACTATAGGTAGCTATGTCTCAGACAAGGCCCT
[G, A]

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AGTAGCTAACAAGCTCTGACTGTCACACAAGGCTTTGTACTGGGAGGCCAGGCTATAGAG
TGGCTCCAGCTTAAAGGGCTGGGAGCTGGGGGACAGTGTCTCAGATTAGGGTCTAACTAG
IT.G]

AAGTTGACTGGAGCTGAGAACAGAGGTTAGGGGCCAAGCAGCAGGGTTGTGGGTCTACTC
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CGGGGTTGGCTGCTAGTCTAAGGGGTGGAGACAAGGACAGAGTTTCAGGTCTGGTCCTTA
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[T,C]

TGCACTACACTTGGGACCACTGCTGCATCATGCCAGGGAGCCTAGAGGTGTCTAAACAGT
TATCCAACAACTGTGATACCCAAGGTTAACTTTCTCTTGTTTTCAGAGGCAGGGAGTACT
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TCCAGGGAACTCCCAGAAATGGTTTGTTTCAGTCAGTTTAGGCTGCTATAAGAGAATATC
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[A,G]

TCCAAGATCAAGGCTCCAGCAGGTTCAGTGTCTGCTGAGTGCTTGTTCTGCTTCGAAGATGGCACCTTTTTGCTGTGTTCTCA

 CTATAAGAGAATATCTTAGAGTGGGTAATCTATCAGCAATAGGAATTTATTGTTCACAAT TCTGGAGGCTGGAAAATCCAAGATCAAGGCTCCAGCAGGTTCAGTGTCTGCTGAGTGCTT [G, C]

TTCTGCTTCGAAGATGGCACCTTTTTGCTGTGTTCTCA